Amendments to the Claims:

This listing of the claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

- 1. (Currently amended) A cytotherapeutic unit suitable for treatment of a patient in need of hematopoietic cells comprising at least about one hundred 1% CD34⁺ cells within a plurality of potent cells, the unit comprising cells from a plurality of sources, wherein said plurality of potent cells comprises isolated CD34⁻, OCT-4⁺ and SSEA3⁻ cells that have been isolated from postpartum placenta perfusate.
 - 2. (Canceled)
 - 3. (Canceled)
 - 4. (Canceled)
- 5. (Previously presented) The cytotherapeutic unit of claim 1 wherein said unit comprises potent cells obtained from fetal cord blood or fetal tissue.
- 6. (Previously presented) The cytotherapeutic unit of claim 1 wherein said unit comprises potent cells obtained from fetal cord blood.
 - 7.-11. (Canceled)
- 12. (Previously presented) The cytotherapeutic unit of claim 1 wherein said potent cells are obtained from at least two individuals.
- 13. (Previously presented) The cytotherapeutic unit of claim 1 wherein said potent cells are obtained from at least five individuals.
 - 14. (Canceled)
- 15. (Original) The cytotherapeutic unit of claim 1 wherein at least one type of cell is excluded from the unit.
- 16. (Original) The cytotherapeutic unit of claim 1 wherein the plurality of potent cells is selected to render the cytotherapeutic unit suitable for therapy for an indicated disease state or condition.
- 17. (Original) The cytotherapeutic unit of claim 16 wherein at least one type of cell is excluded from the unit.
- 18. (Currently amended) A cytotherapeutic unit suitable for the treatment of a patient in need of hematopoietic cells comprising at least two preselected types of potent cells, said unit

comprising cells from a plurality of sources, wherein said potent cells comprise <u>isolated</u> CD34⁻, OCT-4⁺ and SSEA3⁻ cells that have been isolated from postpartum placenta perfusate, and wherein at least about one hundred 1% of said potent cells are CD34⁺.

- 19. (Canceled)
- 20. (Previously presented) The cytotherapeutic unit of claim 18, distributed with a certification of the contents of said cytotherapeutic unit.
- 21. (Previously presented) The cytotherapeutic unit of claim 20 wherein said certification comprises an indication of cells excluded from said cytotherapeutic unit.
- 22. (Previously presented) The cytotherapeutic unit of claim 20 wherein said certification comprises an indication of cells absent from said cytotherapeutic unit.
- 23. (Previously presented) The cytotherapeutic unit of claim 20, wherein said certification indicates how the presence, absence, and/or exclusion of certain cell types render or renders the cytotherapeutic unit suitable for therapy for an indicated disease state or condition.

24.-30.(Canceled)

- 31. (Currently amended) A cytotherapeutic unit suitable for treatment of a patient in need of hematopoietic cells comprising (a) cells obtained from umbilical cord blood and (b) isolated CD34⁻, OCT-4⁺ and SSEA3⁻ cells isolated from postpartum placenta perfusate, wherein at least one type of cell has been removed from the unit, and wherein at least about one hundred 1% of cells remaining in the unit are CD34⁺.
- 32. (Previously presented) The cytotherapeutic unit of claim 31 wherein a plurality of cell types has been removed from the unit.
 - 33. (Canceled)
- 34. (Currently amended) A cytotherapeutic unit suitable for treatment of a patient in need of hematopoietic cells comprising a mixture of cells obtained from umbilical cord blood and <u>isolated</u> CD34⁻, OCT-4⁺ and SSEA3⁻ cells <u>isolated</u> from postpartum placenta perfusate, said <u>mixture of</u> cells comprising a plurality of different types, at least one of the different types having been obtained from a source that differs from a source of another type and wherein at least about one hundred 1% of cells in said cytotherapeutic unit are CD34⁺.
- 35. (Previously presented) The cytotherapeutic unit of claim 34, wherein at least one of said types of cells has been frozen separately from another type of cells.
 - 36. (Original) The cytotherapeutic unit of claim 34, in a frozen state.

- 37. (Previously presented) The cytotherapeutic unit of claim 34, wherein at least one of said cells has been characterized.
 - 38.-53. (Canceled)
- 54. (Previously presented) The cytotherapeutic unit of claim 1, wherein said CD34⁻, OCT-4⁺ and SSEA3⁻ cells are additionally CD10⁺, CD29⁺, CD38⁻, CD44⁺, CD45⁻, CD54⁺, CD90⁺, SH2⁺, SH3⁺, SH4⁺, SSEA4⁻, and ABC-p⁺.
- 55. (Previously presented) The cytotherapeutic unit of claim 18, wherein said cells that are CD34⁻, OCT-4⁺ and SSEA3⁻ cells are additionally CD10⁺, CD29⁺, CD38⁻, CD44⁺, CD45⁻, CD54⁺, CD90⁺, SH2⁺, SH3⁺, SH4⁺, SSEA4⁻, and ABC-p⁺.
- 56. (Previously presented) The cytotherapeutic unit of claim 31, wherein said cells that are CD34⁻, OCT-4⁺ and SSEA3⁻ cells are additionally CD10⁺, CD29⁺, CD38⁻, CD44⁺, CD45⁻, CD54⁺, CD90⁺, SH2⁺, SH3⁺, SH4⁺, SSEA4⁻, and ABC-p⁺.
- 57. (Previously presented) The cytotherapeutic unit of claim 34, wherein said cells that are CD34⁻, OCT-4⁺ and SSEA3⁻ cells are additionally CD10⁺, CD29⁺, CD38⁻, CD44⁺, CD45⁻, CD54⁺, CD90⁺, SH2⁺, SH3⁺, SH4⁺, SSEA4⁻, and ABC-p⁺.